

Use of the correlation matrix approach to define the life detection techniques in a sample curation facility

John Robert Brucato (1), **Andrea Meneghin** (1), Sara Russell (2), Caroline Smith (2), Petra Rettberg (3), Allan Bennet (4), Tom Pottage (4), Aurore Hutzler (5) and the EURO-CARES Team

(1) INAF - Astrophysical Observatory of Arcetri, Firenze, Italy (meneghin@arcteri.astro.it), (2) Natural History Museum, London, UK, (3) Deutsches Zentrum für Luft und Raumfahrt, Cologne, Germany, (4) Public Health England, Salisbury, UK, (5) Naturhistorisches Museum Wien, Vienna, Austria

Abstract

EURO-CARES (European Curation of Astromaterials Returned from Exploration of Space) was a three year (2015-2017), multinational project, funded under the European Commission's Horizon 2020 research programme to develop a roadmap for a European Extra-terrestrial Sample Curation Facility (ESCF). If the samples are brought back to Earth from bodies where there is the possibility of presence of extant or extinct life, there are a wide number of proposed approaches on the techniques to use in order to investigate the presence of biosignatures: [3], [4], [5], etc. All the studies lead to a proposed list of techniques suitable for life detection along with details about the field of application, their efficiency and limits. What is missing is a critical approach able to make a comparison between the techniques in terms of effectiveness, to find a prioritizing ranking. In this paper a quality engineering tool approach, the correlation matrix, was used to support the choice of the techniques for life detection, [1], [2]. The challenge was to analyze and evaluate every technique. To do it, a wide panel of expert was involved. Experts in the following scientific and technological field composed the team: process engineering, mechanical engineering, biology, astrobiology, chemistry. The paper shows how, using a logical flow of analysis, it was possible to identify the critical issues and to highlight the priorities.

1. Introduction

The major drivers we took into account were to define which techniques are really important and which can be considered as optional, rationalize the activity flow inside the curation and provide a support for the design choices of the curation.

Starting from this idea, we focused on the building of a correlation matrix where to correlate the biosignatures with the available techniques. It is known that a number of techniques can detect each biosignature and, at the same time, each technique can be applied for a number of biosignatures. Using the correlation matrix method it is possible to summarize all this information at a glance. It is also possible to give an extra-value to the matrix, trying to be more critical: the idea is not only to determine the correlations between the biosignatures and the techniques, but also to define how strong is each correlation.

2. The correlation matrix

The correlation matrix (Figure 1) shows the correlation between biosignatures and the life detection techniques. According to the matrix approach, the biosignatures were organized per area (morphological, chemical, biochemical, isotopic analysis, and mineralogical), an importance value was given to each techniques, in a range from 1 to 4, and a correlation value was defined, in an exponential range from 0 to 9: 0 if no correlation exists, 1 (low correlation) if the technique is no specific for the biosignature but still usable and/or with medium/low resolution, 3 (medium correlation) if the technique is suitable for the biosignature, although not specific, and/or with medium resolution and 9 (high correlation) if the technique is very specific technique for the biosignature, with high resolution. An extra value was given to disentangle destructive and non-destructive techniques, (1 if the technique is destructive, 1.1 if partially destructive, 1.2 if partially destructive/non-destructive, 1.3 if non-destructive). The numerical results obtained from the correlation matrix are the biosignature occurrence (number of times that the each biosignature is

detected by a different techniques), the techniques occurrences (the number of biosignatures that can be detected by a single techniques), the technique mean value (the technique mean correlation with the detected biosignatures) and finally the technique importance rating calculated, for each column (technique), as the sum of the products of the biosignatures importance, the correlation value and the non-destructive/destructive coefficient.

BIOSIGNATURES		TECHNIQUES																										
Morphological		Size of single cell - Size of targets	Numbers of single bacteria - Number of targets	Population size (colonies)	Chemical composition	Chirality	Organic molecules	DNA, RNA	Organic pigments	Protein	Isotopes, isotopologues	Isotopomers	Elemental analysis	Structure, Mineralogy														
Chemical		4	9	9	4	9	9	9	9	9	9	9	9	9	4	9	9	9	9	9	9	9	9	9	9	9	9	9
Biochemical		4	9	9	4	9	9	9	9	9	9	9	9	9	4	9	9	9	9	9	9	9	9	9	9	9	9	9
Isotopic analysis		4	9	9	4	9	9	9	9	9	9	9	9	9	4	9	9	9	9	9	9	9	9	9	9	9	9	9
Mineralogical		4	9	9	4	9	9	9	9	9	9	9	9	9	4	9	9	9	9	9	9	9	9	9	9	9	9	9
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